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## CONSTRUCTION OF HIV-1/SIVCPZ AMINO ACID ALIGNMENTS

The number of full-length gene sequences is still growing rapidly for all genes. The envelope master alignment now contains 251 full-length sequences. For the purposes of the printed alignments, we have had to limit the number of sequences dramatically. Here we list the criteria we have followed to make the selection.

First, we have decided to end the supremacy of the B clade sequences. More than half (144, to be precise) of the full-length envelope sequences are still subtype B, though the contribution of other subtypes is increasing. We have tried to balance the number of representatives of all subtypes in these alignments. For this, we had to make a heavy selection on subtype B sequences. We have tried to include as many 'classical' sequences as possible. A lot of follow-up work has been done based on lab strains such as HXB2, MN, SF2, and JR-CSF/JR-FL, so these strains are included in the alignments. Although the reference strain for this year's alignments is HXB-2, WEAU is still the reference strain for this Compendium's sister volume, the HIV Immunology Compendium, so this strain is included as well. Furthermore, within subtype B we have tried to represent sequences from diverse geographical origins, so as to represent a broad spectrum of variants. In the case of subtype B, this means that we have included African, Asian and Brazilian variants along with the 'Western' strains.

In a few cases, there were a lot of sequences from non-B subtypes. In these cases we have selected a few representative sequences from each dataset, again with an eye on maintaining geographical diversity. We have left all representatives of group O in the alignment, as these sequences are much more genetically diverse that the subtypes.

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Explanation of Symbols in Alignments		
Symbol	Meaning	
Alignment symbols		
? in consensus	no majority-rule consensus could be determined at this positi	
х	nucleotide missing from codon	
#	frameshift, or codon contains N or illegal character	
\$	stop codon	
Annotation symbols		
	domain boundaries	
/	protein start point	
\	protein end point	
\/	splice site or exon join	
->	start of overlapping coding region	
<-	end of overlapping coding region	
*	cysteine	
^^^ [NxS, NxT]	glycosylation site	
^*^ [NCS, NCT]	glycosylation site with cysteine	
CD4	residue critical for CD4 binding	
cds	coding sequence (indicates regions where two proteins overlap; the overlapping proteins use two different reading frames)	
MHR	major homology region	
nls	nuclear localization signal	
phos site	phosphorylation site	
PKC	protein kinase C binding	
Zn-motif	Zinc finger binding motif	

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Protein	Annotation	Reference
Gag	phos site Ser (111)	Yu, J Biol Chem <b>270</b> :4792 (1995)
Gag	MHR, (284–302)	Otteken, J Virol 70:3407 (1996)
Gag	CyPa (205–241)	Braaten, J Virol <b>70</b> :4220 (1996)
Gag	vpr packaging domain LKSLFG (489–494)	Lu, <i>J Virol</i> <b>69</b> :6873 (1995) Kondo, <i>J Virol</i> <b>70</b> :159 (1996)
Nef	myristylation (1–7)	Huang, J Virol <b>69</b> :93 (1995)
Nef	MHC downmodulation, PK recruitment (26–29)	Piguet, this volume, p. 448 (1999)
Nef	heart of CD4 binding site (55–56)	Piguet, this volume, p. 448 (1999)
Nef	acidic cluster (60–64)	Piguet, this volume, p. 448 (1999)
Nef	(PxxP)3 (67–76)	Huang, J Virol <b>69</b> :93 (1995)
Nef	PKC (75–80)	Huang, J Virol <b>69</b> :93 (1995)
Nef	polypurine tract (89–97)	Huang, J Virol <b>69</b> :93 (1995)
Nef	PAK binding (103–105)	Piguet, this volume, p. 448 (1999)
Nef	Beta turn (128–131)	Huang, J Virol <b>69</b> :93 (1995)
Nef	PxxP (145–148)	Huang, J Virol <b>69</b> :93 (1995)
Nef	COP1 recruitment (152–153)	Piguet, this volume, p. 448 (1999)
Nef	AP recruitment (162–163)	Piguet, this volume, p. 448 (1999)
Nef	V-ATPase and Raf-1 binding (172–173)	Piguet, this volume, p. 448 (1999)
Vpr	alpha helix (16–34)	Cornelissen, ARHR 13:247 (1997)
Vpr	H(S/F)RIG motifs (71–82)	Macreadie, <i>PNAS USA</i> <b>92</b> :2770 (1995)
Vpu	all annotations	Cornelissen, ARHR 13:247 (1997)
Vpr	LR domain (60–82)	Wang, Gene 178:7 (1996)